CYCLIC BIOISOSTERES OF DERIVATIVES OF A PURINE SYSTEM AND COMPOSITION BASED THEREON

Abstract

The invention relates to cyclic bioisosteres of derivatives of a purine system having a general structural formula

$$R^{1}$$
 R^{1}
 R^{1}
 R^{1}
 R^{1}
 R^{1}
 R^{2}
 R^{1}
 R^{2}
 R^{2}
 R^{3}
 R^{4}
 R^{4

$$R^{1} = -H, -NH_{2}, -Br, -C1, -OH, -COOH,$$

$$A = -N = \text{ at } B = -N = , Z = -CH - ,$$

$$A = -CH = at B = -N = Z = -CH - Z$$

$$A = -CH = at B = -N = Z = -N = Z$$

$$A = -CH = at B = -CH = Z = -CH = Z$$

$$A = -CH = at B = -CH = Z = -N = A$$

and their pharmacologically acceptable salts having a normalizing effect on endocellular processes, in particular, it is capable eliminating endocellular metabolic acidosis and capable of binding excessively formed free radicals, in particular, free-radical forms of oxygen, capable of normalizing the nitrergic mechanisms of the cells, and also capable of interreacting with adenosine-sensitive receptors on the membrane of non-nuclear cells and in nuclei-containing cells to decrease the aggregation of thrombocytes. The compounds according to the invention have hepatoprotective effect and can be used for producing pharmaceutical compositions on their base.